

EXHIBIT A

Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations

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Product Details for ANDA 210531

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DIMETHYL FUMARATE (DIMETHYL FUMARATE)_
120MG
Marketing Status: Prescription

Active Ingredient: DIMETHYL FUMARATE
Proprietary Name: DIMETHYL FUMARATE
Dosage Form; Route of Administration: CAPSULE, DELAYED RELEASE; ORAL
Strength: 120MG
Reference Listed Drug: No
Reference Standard: No
TE Code: AB
Application Number: A210531
Product Number: 001
Approval Date: Aug 17, 2020
Applicant Holder Full Name: MYLAN PHARMACEUTICALS INC
Marketing Status: Prescription
Patent and Exclusivity Information (patent_info.cfm?
Product_No=001&Appl_No=210531&Appl_type=A)

DIMETHYL FUMARATE (DIMETHYL FUMARATE)_
240MG

Marketing Status: Prescription

Active Ingredient: DIMETHYL FUMARATE

Proprietary Name: DIMETHYL FUMARATE

Dosage Form; Route of Administration: CAPSULE, DELAYED RELEASE; ORAL

Strength: 240MG

Reference Listed Drug: No

Reference Standard: No

TE Code: AB

Application Number: A210531

Product Number: 002

Approval Date: Aug 17, 2020

Applicant Holder Full Name: MYLAN PHARMACEUTICALS INC

Marketing Status: Prescription

Patent and Exclusivity Information (patent_info.cfm?

Product_No=002&Appl_No=210531&Appl_type=A)